(g) * * *

(12) Have failed to find employment after utilizing services provided under title I of the Workforce Innovation and Opportunity Act;

(13) Are homeless or at risk for homelessness; or

(14) Are formerly incarcerated individuals as defined in § 641.140.

Angela Hanks,

Acting Assistant Secretary for Employment and Training, Labor.

[FR Doc. 2022–02681 Filed 2–11–22; 8:45 am]

BILLING CODE 4510–FN–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 73

[Docket No. FDA–2022–C–0098]

Motif FoodWorks, Inc.; Filing of Color Additive Petition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notification of petition.

SUMMARY: The Food and Drug Administration (FDA or we) is announcing that we have filed a petition, submitted by Motif FoodWorks, Inc., proposing that the color additive regulations be amended to provide for the safe use of myoglobin as a color additive in meat and poultry analogue products.

DATES: The color additive petition was filed on December 13, 2021.

ADDRESSES: For access to the docket to read background documents or comments received, go to *https://www.regulations.gov* and insert the docket number found in brackets in the heading of this document into the ''Search'' box and follow the prompts, and/or go to the Dockets Management Staff, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Ellen Anderson, Center for Food Safety and Applied Nutrition, Food and Drug Administration, 5001 Campus Dr., College Park, MD 20740, 240–402–1309.

SUPPLEMENTARY INFORMATION: Under section 721(d)(1) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 379e(d)(1)), we are giving notice that we have filed a color additive petition (CAP 2C0322), submitted by Motif FoodWorks, Inc., 27 Drydock Ave., 2nd Floor, Boston, MA 02210. The petition proposes to amend the color additive regulations in part 73 (21 CFR part 73), ''Listing of Color Additives Exempt from Certification,'' to provide for the safe use of myoglobin as a color additive in meat and poultry analogue products.

The petitioner has claimed that this action is categorically excluded under 21 CFR 25.32(r) because the substance occurs naturally in the environment, and the action does not alter significantly the concentration or distribution of the substance, its metabolites, or degradation products in the environment. In addition, the petitioner has stated that, to their knowledge, no extraordinary circumstances exist that would warrant at least an environmental assessment (see 21 CFR 25.21). If FDA determines a categorical exclusion applies, neither an environmental assessment nor an environmental impact statement is required. If FDA determines a categorical exclusion does not apply, we will request an environmental assessment and make it available for public inspection.

Dated: February 7, 2022.

Lauren K. Roth,

Associate Commissioner for Policy.

[FR Doc. 2022–03109 Filed 2–11–22; 8:45 am]

BILLING CODE 4164–01–P

ENVIRONMENTAL PROTECTION AGENCY

40 CFR Part 52

[EPA–R10–OAR–2020–0684, FRL–9402–01–R10]

Air Plan Approval; OR; Air Contaminant Discharge Permit Fee Revision

AGENCY: Environmental Protection Agency (EPA).

ACTION: Proposed rule.

SUMMARY: The Environmental Protection Agency (EPA) is proposing to approve a revision to the Oregon State Implementation Plan (SIP) submitted on November 5, 2020. The revision establishes new fees to be paid by stationary sources of air contaminants submitting notices of intent to construct. The revision also adds a new basic air contaminant discharge permit category to allow certain minor sources, that would otherwise be required to obtain a general, simple, or standard permit, the option to qualify for a basic permit.

DATES: Comments must be received on or before March 16, 2022.

ADDRESSES: Submit your comments, identified by Docket ID No. EPA–R10–OAR–2020–0684, at *https://www.regulations.gov.* Follow the online instructions for submitting comments. Once submitted, comments cannot be edited or removed from *https://www.regulations.gov.* The EPA may publish any comment received to its public docket. Do not electronically submit any information you consider to be Confidential Business Information (CBI) or other information the disclosure of which is restricted by statute. Multimedia submissions (audio, video, etc.) must be accompanied by a written comment. The written comment is considered the official comment and should include discussion of all points you wish to make. The EPA will generally not consider comments or comment contents located outside of the primary submission (*i.e.,* on the web, cloud, or other file sharing system). For additional submission methods, the full EPA public comment policy, information about CBI or multimedia submissions, and general guidance on making effective comments, please visit *https://www.epa.gov/dockets/commenting-epa-dockets.*

FOR FURTHER INFORMATION CONTACT: Kristin Hall, EPA Region 10, 1200 Sixth Avenue, Suite 155, Seattle, WA 98101, at (206) 553–6357 or *hall.kristin@epa.gov.*

SUPPLEMENTARY INFORMATION: Throughout this document, wherever ''we'' or ''our'' is used, it is intended to refer to the EPA.

Table of Contents

I. Background

A. State Implementation Plan

Each state has a State Implementation Plan (SIP) containing the control measures and strategies used to attain and maintain the national ambient air quality standards (NAAQS) established by the EPA for the criteria pollutants (carbon monoxide, lead, nitrogen dioxide, ozone, particulate matter, sulfur dioxide). Section 110 of the Clean Air Act spells out the requirements for each SIP, including but not limited to air pollution control regulations, emissions inventories, ambient air monitoring, enforcement mechanisms, and authority to revise the SIP as needed.

Revisions to the SIP are adopted by the state and submitted to the EPA for review. The EPA approves and codifies